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10/540,814	01/09/2006	Kenji Miyazaki	18962	6341
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Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No.	Applicant(s)	
	10/540,814	MIYAZAKI ET AL.	
	Examiner	Art Unit	
	ROBERT XU	1797	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) Responsive to communication(s) filed on 09 January 2006.
 2a) This action is FINAL. 2b) This action is non-final.
 3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) Claim(s) 1-17 is/are pending in the application.
 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
 5) Claim(s) _____ is/are allowed.
 6) Claim(s) _____ is/are rejected.
 7) Claim(s) _____ is/are objected to.
 8) Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) The specification is objected to by the Examiner.
 10) The drawing(s) filed on 24 June 2005 is/are: a) accepted or b) objected to by the Examiner.
 Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
 Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
 11) The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
 a) All b) Some * c) None of:
 1. Certified copies of the priority documents have been received.
 2. Certified copies of the priority documents have been received in Application No. _____.
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|---|---|
| 1) <input checked="" type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ . |
| 3) <input checked="" type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08)
Paper No(s)/Mail Date <u>6/24/2005, 9/15/2006</u> . | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

1. An examiner's amendment to the record appears below. Should the changes and/or additions be unacceptable to applicant, an amendment may be filed as provided by 37 CFR 1.312. To ensure consideration of such an amendment, it MUST be submitted no later than the payment of the issue fee.

Authorization for this examiner's amendment was given in a telephone interview with Xiaochun Zhu, Esq., Ph.D. on 10/06/08.

The application has been amended as follows:

In claim 4: delete [any of] and [to 3] and replace [Claims] with -- Claim --;

In claim 5: delete [any of] and [to 4] and replace [Claims] with -- Claim --;

Claims 1-17 are pending and have been fully considered.

Claim Rejections - 35 USC § 112

2. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

3. Claim 1 is rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 1, page 322, lines 8-10 recites "calculating the relative peak intensity on the basis of said peak intensity of highest cationic species peak". It is not clear, weather the "highest cationic species peak" is the one from the group of cationic species peaks corresponding to the third group of anionic species peaks (the highest in the group), or the one from the first group of cationic species peaks (the highest in all groups).

Claim 1, page 322, lines 12-14 recites "calculating the relative peak intensity on the basis of said peak intensity of highest anionic species peak". It is not clear, weather the "highest anionic species peak" is the one from the third group of anionic species peaks (the highest in the third group), or the one from the first group of anionic species peaks (the highest in all groups).

Claim Rejections - 35 USC § 103

4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

5. The factual inquiries set forth in *Graham v. John Deere Co.*, 383 U.S. 1, 148 USPQ 459 (1966), that are applied for establishing a background for determining obviousness under 35 U.S.C. 103(a) are summarized as follows:

1. Determining the scope and contents of the prior art.
2. Ascertaining the differences between the prior art and the claims at issue.
3. Resolving the level of ordinary skill in the pertinent art.
4. Considering objective evidence present in the application indicating obviousness or nonobviousness.

6. **Claims 1, 2 and 4-6** are rejected under 35 U.S.C. 103(a) as being unpatentable over Tsugita et al. (Electrophoresis, 1998) (Tsugita) in view of Covey et al. (US Patent 5,952,653) (Convey) and Xu et al. (Analytical Biochemistry, 1997) (Xu).

In regard to Claim 1, Tsugita teaches a method of analyzing C-terminal sequence of a peptide by means of mass spectrometry.

Tsugita teaches pretreatment step using acetic anhydride react with the dried peptide to acetylate the N-terminus and form an oxazolone at the C-terminal carboxyl group (see page 930, right col. 3rd paragraph). While Tsugita does not specifically teach acetylation of the side chain of lysine residue, acetylation of the side chain of lysine residue is the inherent result of reacting acetic anhydride with peptide that has lysine residue.

Tsugita teaches a step of allowing acetic anhydride to act on the dry peptide in the presence of pentafluoropropionic methyl ester (PFPMMe) to successively release the C-terminal amino acid by the cleavage of the 5-oxazolone ring (see page 930, right col. 3rd paragraph). PFPMMe is similar to pentafluoropropionic acid (PFPA) in structure and reaction. Tusgita also teaches using PFPA in the cleavage of predetermined position of

the peptide (see page 932, left col. 4th paragraph). It would have been obvious to ordinary skill in the art to substitute PFPMe with PFPA in the step of releasing the C-terminal amino acids in Tsugita method, because these compounds have similar structures and similar effects.

Tsugita teaches hydrolysis treatment step by allowing water molecules to act on the C-terminal-deleted peptides in the presence of dimethylamino ethanol (DMAE) (see page 930, right col. 3rd paragraph).

Tsugita teaches measuring the decreases in molecular weight associated with the successive release of the C-terminal amino acids by mass spectra (see Table 3).

Tsugita does not specifically teach allowing trypsin to act on the sample to cleave peptide for mass spectrometer analysis. Tsugita teaches that highly specific proteases have been used for specific fragmentation in the peptide-mass fingerprinting technique (see page 931, left col. 2nd paragraph). Trypsin is one of highly specific proteases commonly used. Covey teaches using trypsin to cleave peptide for mass spectrometry analysis (see abstract). Covey further teaches that the tryptic fragments ions are predominantly doubly charge cationic species, because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not containing an arginine or a lysine is single charged (see col. 5, lines 58-60). At the time of the invention, it would have been obvious to one of ordinary skill in the art to use trypsin to cleave the peptide before mass spectrometry analysis, as taught by Covey, in the method of Tsugita, because smaller peptides are easier to analyze by mass spectrometry.

Tsugita does not teach the protocol of analyzing mass spectra by comparing the peaks of cationic species with the peaks of the anionic species as described in the steps 1-9 in the instant claim. As has been discussed above, Covey teaches that tryptic fragments ions are predominantly doubly charge cationic species because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not containing an arginine or a lysine is single charged (see col. 5, lines 58-60). The method

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of analyzing mass spectra according to the expected charges of the species is well known in the art.

Xu teaches that fragmentation patterns in positive mode and negative mode are complementary for the elucidation of the peptide chain sequence (see abstract). Xu further teaches correlating peaks that only differ in a chemical group, e.g. OH (mass of 18) or N-acetyl glucosamine (MurNAC mass of 203) because of the reaction (see page 10, right col.). Xu's teaching is similar to the criteria 5a-1, 5a-2, 5a-3, 5b-1, 5b-2, 5b-3 of the instant claim.

Applicant is advised that the rationale to support a conclusion that the claim would have been obvious is that all the claimed elements were known in the prior art and one skilled in the art could have combined the elements as claimed by known methods with no change in their respective functions, and the combination yielded nothing more than predictable results to one of ordinary skill in the art. (see KSR, 550 U.S. at ___, 82 USPQ2d at 1395) (see MPEP 2143). In that regard, one of ordinary skill in the art could have utilized the charge difference between normal tryptic fragments and C-terminal tryptic fragment as taught by Covey, and correlated the fragmentation patterns in positive mode and negative mode in relative intensity and further correlated with the special group difference due to chemical reaction, as taught by Xu to elucidate the peptide chain sequence by calculating the mass difference between the C-terminal successive deleted peptides as taught by Tsugita, with the predictable result.

In regard to Claim 2, it is well known in the art that spiked noise in mass spectra usually has narrower full-width of half maximum than normal peak of signal. Therefore, removing peaks of spiked noise based on full-width of half maximum would have been obvious to ordinary skill in the art. The peak smoothing and smoothing algorithms are also well known in the art. Therefore, peak smoothing would have been obvious to ordinary skill in the art.

In regard to Claim 4, the normal tryptic fragments carry two positive charges as taught by Covey. Under the same reason, when C-terminal fragment has arginine at the C-terminal (CFAC), it will also carry two positive charges. However, just like other C-terminal fragments, CFAC will also have an adjacent fragment peak that has molecular

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weight difference from CFAC equals formula weight of natural chain amino acid or acylated amino acid as taught by Tsugita. It would have been obvious to ordinary skill in the art to use the criteria of adjacent peaks to judge if the strong cationic peak is CFAC based on teaching of Tsugita and Convey.

In regard to Claim 5, Tsugita teaches measuring the decrease in molecular weight associated with successive release of the C-terminal amino acids (see Table 3). Tsugita teaches using MALDI-TOF-MS for the measurement (see page 931, left col. 2nd paragraph). Xu teaches considering fragmentation patterns in both cationic species (positive mode) and anionic species (negative modes) (see abstract). At the time of the invention, it would have been obvious for ordinary skill in the art to consider both cationic species and anionic species as taught by Xu in Tsugita's method to obtain the peptide sequence, because Xu specifically teaches that the fragmentation in positive mode and negative mode are complementary for elucidation of the peptide sequence.

In regard to Claim 6, Tsugita teaches a process for releasing the C-terminal amino acids successively. Tsugita teaches pretreatment step using acetic anhydride and acetic acid vapor react with the dried peptide at 60°C to acetylate the N-terminus and form an oxazolone at the C-terminal carboxyl group (see page 930, right col. 3rd paragraph). Tsugita does not specifically teach acetylation of the side chain of lysine residue. Acetylation of the side chain of lysine residue is the inherent result of reacting acetic anhydride with peptide that has lysine residue.

Tsugita teaches a step of allowing acetic anhydride vapor to act on the dry peptide in the presence of pentafluoropropionic methyl ester (PFPMe) vapor to successively release the C-terminal amino acid by the cleavage of the 5-oxazolone ring at 5°C (see page 930, right col. 3rd paragraph). PFPMe is similar to pentafluoropropionic acid (PFPA) in structure and reaction. Tsugita also teaches using PFPA in the cleavage of predetermined position of the peptide (see page 932, left col. 4th paragraph). It would have been obvious to ordinary skill in the art to substitute PFPMe with PFPA in the step of releasing the C-terminal amino acids. Tsugita allows the reaction at 5°C. The court has held that [W]here the general conditions of a claim are disclosed in the prior art, it is not inventive to discover the optimum or workable ranges by routine experimentation (*In*

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re Aller, 220 F.2d 454, 456, 105 USPQ 233, 235 (CCPA 1955)). In that regard, although 5°C is outside the range of 15-60°C in the instant claim, it would have been obvious for ordinary skill in the art to optimize the reaction temperature by routine experimentation.

Tsugita does not literally teach removing the remaining alkanoic acid anhydride and perfluoroalkanoic acid in a dry state at the end of C-terminal cleaving reaction. However, this removing step at the end of the reaction is inherent part of the reaction step, because when the reaction is complete, the remaining reagents should be removed. Tsugita teaches hydrolysis treatment step by allowing water molecules to act on the C-terminal-deleted peptides in the presence of dimethylamino ethanol (DMAE). (see page 930, right col. 3rd paragraph). Again, the removing remaining basic nitrogen-containing organic is inherent part of the reaction step.

Tsugita does not specifically teach allowing trypsin to act on the sample to cleave peptide for mass spectrometer analysis. Tsugita teaches that highly specific proteases have been used for specific fragmentation in the peptide-mass fingerprinting technique (see page 931, left col. 2nd paragraph). Trypsin is one of highly specific proteases commonly used. Covey teaches using trypsin to cleave peptide for mass spectrometry analysis (see abstract). Covey further teaches that the tryptic fragments ions are predominantly doubly charge cationic species because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not contain an arginine or a lysine is signally charge (see col. 5, lines 58-60). Since the side chain of lysine is protected by acetylation treatment as discussed above, only arginine site will be cut by trypsin. At the time of the invention, it would have been obvious to one of ordinary skill in the art to use trypsin to cleave the peptide before mass spectrometry analysis as taught by Covey in the method of Tsugita, because smaller peptides are easier to analyze by mass spectrometry.

Removing trypsin at the end of the reaction is inherent part of the reaction. Desalting treatment is commonly used for changing buffer or removing buffer solution component. It would have been obvious to ordinary skill in he art to use desalting

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treatment to remove the trypsin from the buffer solution to stop the reaction. Drying sample is a required step before performing MALDI-TOF-MS.

Tsugita teaches using MALDI-TOF-MS for measuring the molecular weight of the peptide fragments (see page 931, left col. 2nd paragraph). Tsugita does not teach analyzing mass spectra by comparing the peaks of cationic species with the peaks of the anionic species. As has been discussed above, Covey teaches that tryptic fragments ions are predominantly doubly charge cationic species because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not containing an arginine or a lysine is signally charge (see col. 5, lines 58-60). The method of analyzing mass spectra according to the expected charges of the species is well known in the art. Xu teaches that fragmentation patterns in positive mode and negative mode are complementary for the elucidation of the peptide chain sequence (see abstract). It would have been obvious to one of ordinary skill in the art to utilize the charge difference between normal tryptic fragments and C-terminal tryptic fragment as taught by Covey and correlate the fragmentation patterns in positive mode and negative mode as taught by Xu to elucidate the peptide chain sequence by means of MALDI-TOF-MS as taught by Tsugita.

7. **Claim 3 is rejected under 35 U.S.C. 103(a) as being unpatentable over Tsugita in view of Covey and Xu, as applied to claims 1-2 and 4-6 above, and further in view of Harris et al. (Rapid Communications in mass spectrometry, 2002) (Harris).**

In regard to Claim 3, Tsugita in view of Convey and Xu do not teach using the fragments of trypsin autolysis as the internal standard to calibrate mass spectra. The molecular weights and charges of peptide fragments derived from the trypsin autolysis are well known in the art. Harris teaches using trypsin autolysis fragments as mass calibrants in matrix-assisted laser desorption/ionization time-of-flight mass spectrometry (see abstract). Therefore, it would have been obvious to ordinary skill in the art to modify Tsugita-Convey-Xu's method by calibrating mass spectra using the peptide fragments of trypsin autolysis, as taught by Harris.

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8. **Claims 7-17** are rejected under 35 U.S.C. 103(a) as being unpatentable over Tsugita in view of Covey and Xu as applied to 1,2 and 4-6 above, and further in view of Vogt et al. (Polymer Bulletin, 1996) (Vogt)

In regard to Claim 7, Tsugita teaches separating target peptide by electrophoresis before process for C-terminal analysis (see page 929, right col. 2nd paragraph). Tsugita in view of Convey and Xu do not teach releasing the C-terminal amino acids successively while the peptide is bound on a gel carrier. Tsugita teaches that the pretreatment and cleavage sub-step of the procedure needs to be carried out in the absence of water (page 931, right col. 2nd paragraph, last 2 lines; page 930, 3rd paragraph). Therefore, the target protein has to be extracted from the gel and then dried to remove water or electroblotted to an Immobilon-CD membrane. The consequence of extracting peptide from a gel after separation is that some peptide will be lost during the process (it is true to any peptide or protein processing). Therefore, sequencing the peptide while it is bound to a gel would avoid the loss of peptide during extraction.

Tsugita does not teach a step of using polar aprotic solvent to remove water. However, using polar aprotic solvent to remove water is known in the art.

Tsugita teaches applying N-acylation protection by alkanoic acid anhydride at 60°C (see page 930, right col. 3rd paragraph). Tsugita does not teach using dipolar-aprotic solvent to swollen the gel so that the pretreatment and C-terminal cleavage reaction could be carried out on the protein bound to the original gel right after electrophoresis. Vogt teaches a new non-aqueous swelling system; specifically he teaches that carboxymethyl cellulose (CMC) gel treated with a dipolar aprotic solvent like *N,N*-dimethylacetamide with *p*-toluenesulfonic acid yields a high reactive gel-suspension of the polymer (see abstract). This dipolar aprotic solvent can remove water from the swollen gel in one step (see page 550, 3rd paragraph), thus allowing a direct esterification of the hydroxyl group of CMC (see abstract). At the time of the invention, it would have been obvious to one of ordinary skill in the art to use polar aprotic solvent to remove water and use dipolar aprotic solvent to remove water from the gel carrier bound with the target protein, as taught by Vogt with reasonable expectation that this

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would allow Tsugita-Convey-Xu's procedure to be carried out on the target protein kept on the gel carrier. Sequencing the peptide while it is bound to a gel would avoid the loss of peptide during extraction. Also because Vogt specifically teaches that dipolar aprotic solvent can remove water from gel while keep the gel swollen.

Tsugita teaches a step of allowing acetic anhydride to act on the dry peptide in the presence of pentafluoropropionic methyl ester (PFPM_e) to successively release the C-terminal amino acid by the cleavage of the 5-oxazolone ring (see page 930, right col. 3rd paragraph). PFPM_e is similar to pentafluoropropionic acid (PFPA) in structure and reaction. Tsugita also teaches using PFPA in the cleavage of predetermined position of the peptide (see page 932, left col. 4th paragraph). It would have been obvious to ordinary skill in the art to substitute PFPM_e with PFPA in the step of releasing the C-terminal amino acids in Tsugita method, because these compounds have similar structures and similar effects.

Tsugita does not teach using polar aprotic solvent to remove dipolar aprotic solvent. Polar aprotic solvent and dipolar aprotic solvent are both aprotic solvents and very similar. It is well known that they can be used to remove each other.

Tsugita teaches hydrolysis treatment with an aqueous solution of tertiary amine compound (DMAE) after successive release of C-terminal amino acids (see page 930, right col. 3rd paragraph).

Tsugita does not specifically teach allowing trypsin to act on the sample to cleave peptide for mass spectrometer analysis. Tsugita teaches that highly specific proteases have been used for specific fragmentation in the peptide-mass fingerprinting technique (see page 931, left col. 2nd paragraph). Trypsin is one of highly specific proteases commonly used. Covey teaches using trypsin to cleave peptide for mass spectrometry analysis (see abstract). Covey further teaches that the tryptic fragments ions are predominantly doubly charge cationic species because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not contain an arginine or a lysine is signally charge (see col. 5, lines 58-60). Since the side chain of lysine is protected by acetylation treatment, only arginine site will be cut by trypsin. At the time of

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the invention, it would have been obvious to one of ordinary skill in the art to use trypsin to cleave the peptide before mass spectrometry analysis as taught by Covey in the method of Tsugita, because smaller peptides are easier to analyze by mass spectrometry.

Desalting treatment is commonly used for changing buffer or removing buffer solution component. It would have been obvious to ordinary skill in he art to use desalting treatment to remove the trypsin from the buffer solution to stop the reaction. Drying sample is a required step before performing MALDI-TOF-MS.

Tsugita teaches using MALDI-TOF-MS for measuring the molecular weight of the peptide fragments (see page 931, left col. 2nd paragraph). Tsugita does not teach analyzing mass spectra by comparing the peaks of cationic species with the peaks of the anionic species. As has been discussed above, Covey teaches that tryptic fragments ions are predominantly doubly charge cationic species because arginine and lysine are both very basic and each picks up a positive charged proton in solution (see col. 2. lines 63-67; Col. 3, lines 39-42), and the C-terminal tryptic fragment not containing an arginine or a lysine is signally charge (see col. 5, lines 58-60). The method of analyzing mass spectra according to the expected charges of the species is well known in the art. Xu teaches that fragmentation patterns in positive mode and negative mode are complementary for the elucidation of the peptide chain sequence (see abstract). It would have been obvious to one of ordinary skill in the art to utilize the charge difference between normal tryptic fragments and C-terminal tryptic fragment as taught by Covey and correlate the fragmentation patterns in positive mode and negative mode as taught by Xu to elucidate the peptide chain sequence by means of MALDI-TOF-MS as taught by Tsugita.

In regard to Claims 8 and 9, Tsugita teaches that 20% acetic anhydride is used in the first step of the procedure for applying N-acetylation protection to the N-terminal of the protein and for forming oxazolone at C-terminal of the protein and 5% PFPMe is used in the second step to react with oxazolone (page 930, right col. 3rd paragraph). Tsugita does not specifically teach maintaining acetic anhydride in the second step. However, since the function of acetic anhydride is to form oxazolone at C-terminal for

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perfluoroalkanoic acid to act on in the second step, it would have been obvious to one of ordinary skill in the art to recognize that maintaining the concentration of acetic anhydride in the second step may benefit the reaction. Therefore, modified method of Tsugita teaches using symmetric anhydride of linear -chain alkanoic acid having 2 carbons (acetic anhydride) for the formation of 5-oxazolone and subsequently release of C-terminal amino acids in association with cleavage of the 5-oxazolone ring (see page 930, right col. 3rd paragraph). Asymmetric anhydride has similar structure and function as symmetric anhydride. At time of the invention it would have been obvious to one of ordinary skill in the art to substitute symmetric anhydride with asymmetric anhydride with reasonable expectation that the asymmetric anhydride would also cleave the 5-oxazolone ring.

In regard to Claim 10, Tsugita teaches using acetic anhydride as the alkanoic acid anhydride (see page 930, right col. 3rd paragraph).

In regard to Claim 11, the pKa of PFPA is in a range of 0.3 to 2.5.

In regard to Claim 12, PFPA has 3 carbon atoms.

In regard to Claim 13, modified method of Tsugita teaches that the ratio of PFPA (5%) to acetic anhydride (20%) would be 1:4 (see page 930, right col. 3rd paragraph). In the instant Claim, the lower limit of the ratio is 20:100 or 1:5. Applicant is advised that generally, differences in concentration or temperature will not support the patentability of subject matter encompassed by the prior art unless there is evidence indicating such concentration or temperature is critical. "[W]here the general conditions of a claim are disclosed in the prior art, it is not inventive to discover the optimum or workable ranges by routine experimentation." *In re Aller*, 220 F.2d 454, 456, 105 USPQ 233, 235 (CCPA 1955). Therefore, it would have been obvious to one of ordinary skill in the art to discover the optimum ratio of perfluoroalkanoic acid to alkanoic acid anhydride by routine experimentation.

In regard to Claims 14-16, Tsugita teaches that acetic anhydride is used in the pretreatment step of applying N-acylation protection (see page 930, right col. 3rd paragraph).

In regard to Claim 17, Tsugita teaches using acetic anhydride as the alkanoic acid anhydride in the pretreatment step of applying N-acylation protection. Modified method of Tsugita teaches using acetic anhydride in combination with PFPA for the formation of 5-oxazolone and subsequent release of C-terminal amino acids in association with the cleavage of the 5-oxazolone ring (see page 930, right col. 3rd paragraph).

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to ROBERT XU whose telephone number is (571)270-5560. The examiner can normally be reached on Mon-Thur 7:30am-5:00pm, Fri 7:30am-4:00pm.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Jill Warden can be reached on (571)272-1267. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300. Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

10/17/2008

/Yelena G. Gakh/
Primary Examiner, Art Unit 1797

RX